Project Title	Funding	Institution	
Understanding the Genetic Architecture of Rett Syndrome - an Autism Spectrum Disorder	\$30,000	Cold Spring Harbor Laboratory	
Undergraduate Research Award	\$3,000	Texas A&M University	
TrkB agonist therapy for sensorimotor dysfunction in Rett syndrome	\$5,867	Case Western Reserve University	
Translational Regulation of Adult Neural Stem Cells	\$372,633	University of Wisconsin	
Translational dysregulation in autism pathogenesis and therapy	\$250,000	Massachusetts General Hospital	
Translation, Synchrony, and Cognition	\$380,953	New York University	
The role of UBE3A in autism: Is there a critical window for social development?	\$54,450	Erasmus University Medical Center	
The role of Shank3 in neocortex versus striatum and the pathophysiology of autism	\$0	Duke University	
THE ROLE OF MECP2 IN RETT SYNDROME	\$356,699	University of California, Davis	
The Role of Glia in Fragile X Syndrome	\$0	Johns Hopkins University	
Thalamocortical circuit defects in developmental brain disorders	\$490,462	University of Maryland	
Tet-mediated Epigenetic Modulation in Autism	\$603,129	Emory University	
Testing the ribosomal protein S6 as treatment target and biomarker in autism spectrum disorders	\$0	Cincinnati Children's Hospital	
Targeting the PI3K Enhancer PIKE to Reverse FXS-associated Phenotypes	\$160,000	Emory University	
Synaptic Phenotype, Development, and Plasticity in the Fragile X Mouse	\$395,642	MICHIGAN STATE UNIVERSITY	
Supplement to The Emergence and Stability of Autism in Fragile X Syndrome	\$82,061	UNIVERSITY OF SOUTH CAROLINA	
Role of UBE3A in the Central Nervous System	\$321,269	University of North Carolina	
Role of Serotonin Signaling during Neural Circuitry Formation in Autism Spectrum Disorders	\$0	Massachusetts Institute of Technology	
Role of MEF2 and neural activity in cortical synaptic weakening and elimination	\$388,354	UT SOUTHWESTERN MEDICAL CENTER	
Role of GABA interneurons in a genetic model of autism	\$0	Yale University	
Rescuing synaptic and circuit deficits in an Angelman syndrome mouse model	\$60,000	Arizona Board of Regents, University of Arizona	
Rapid screening for cortical circuit dysfunction in autism-related mouse models	\$0	University of California, Berkeley	
Profiles and Predictors of Pragmatic Language Impairments in the FMR1 Premutation	\$55,796	UNIVERSITY OF SOUTH CAROLINA	
Probing the neural basis of social behavior in mice	\$0	Massachusetts Institute of Technology	
Probing the Molecular Mechanisms Underlying Autism: Examination of Dysregulated Protein Synthesis	\$0	National Institutes of Health	
Probing synaptic receptor composition in mouse models of autism	\$124,998	Boston Children's Hospital	
Presynaptic Fragile X Proteins	\$249,000	DREXEL UNIVERSITY	
Pragmatic language and social-emotional processing in autism, fragile X, and the FMR1 premutation	\$0	Northwestern University	

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PPAR/SIRT1 PATHWAY IN C. ELEGANS	\$22,740	Children's Hospital of Philadelphia	
Potassium channels as therapeutic targets in autism	\$60,000	Administrators of the Tulane Educational Fund	
Novel candidate mechanisms of fragile X syndrome	\$248,235	UNIVERSITY OF MICHIGAN	
New Models For Astrocyte Function in Genetic Mouse Models of Autism Spectrum Diso	\$396,250	CLEVELAND CLINIC LERNER COM-CWRU	
Neurotrophic Factor Regulation of Gene Expression	\$618,134	Harvard University	
Neuropathology of the Shank3 mouse model for autism	\$0	University of Louisville	
Neuronal translation in Tsc2+/- and Fmr1-/y mutant ASD mouse models	\$62,500	The Trustees of Columbia University in the City of New York	
Neuronal Activity-Dependent Regulation of MeCP2	\$600,383	Harvard University	
Neurobiology of Rai1, a critical gene for syndromic ASDs	\$87,500	The Board of Trustees of the Leland Stanford Junior University (Stanford)	
Neurobiological Mechanism of 15q11-13 Duplication Autism Spectrum Disorder	\$380,625	BETH ISRAEL DEACONESS MEDICAL CENTER	
Neural mechanisms underlying autism behaviors in SCN1A mutant mice	\$100,000	University of Washington	
Neural Correlates of the Y Chromosome in Autism: XYY Syndrome as a Genetic Model	\$0	Children's Hospital of Philadelphia	
Neural Correlates of the Y Chromosome in Autism: XYY Syndrome as a Genetic Model	\$0	Nemours Children's Health System, Jacksonville	
Neural and cognitive discoordination in autism-related mouse models	\$280,480	New York University	
Multigenic basis for autism linked to 22q13 chromosomal region	\$125,000	Hunter College of the City University of New York (CUNY) jointly with Research Foundation of CUNY	
mTOR modulation of myelination	\$179,659	Vanderbilt University	
MRI Biomarkers of Patients with Tuberous Sclerosis Complex and Autism	\$727,821	CHILDREN'S HOSPITAL CORPORATION	
Mouse Model of Dup15q Syndrome	\$32,635	Texas AgriLife Research	
Motor cortex plasticity in MeCP2 duplication syndrome	\$30,000	Baylor College of Medicine	
Modeling Pitt-Hopkins Syndrome, an Autism Spectrum Disorder, in Transgenic Mice Harboring a Pathogenic Dominant Negative Mutation in TCF4	\$0	University of North Carolina	
Modeling Microglial Involvement in Autism Spectrum Disorders, with Human Neuro-glial Co-cultures	\$30,000	Whitehead Institute for Biomedical Research	
Mechanisms underlying word learning in fragile X syndrome and nonsyndromic ASD	\$156,333	University of California, Davis	
Mechanisms underlying the Cerebellar Contribution to Autism in Mouse Models of Tuberous Sclerosis Complex	\$190,458	UT SOUTHWESTERN MEDICAL CENTER	
Mechanisms of synapse elimination by autism-linked genes	\$0	University of Texas Southwestern Medical Center	
Mechanisms of Motor Skill Learning in the Fragile X Mouse Model	\$300,434	University of Nebraska	
Mechanisms of mGluR5 function and dysfunction in mouse autism models	\$410,720	UT SOUTHWESTERN MEDICAL CENTER	
Mechanisms and Rescue of Neural Circuit Dysfunction in Mecp2 Mutant Mice	\$92,578	Baylor College of Medicine	

Project Title	Funding	Institution	
Mapping the Neurobehavioral Phenotype in Phelan McDermid Syndrome	\$0	ICAHN SCHOOL OF MEDICINE AT MOUNT SINAI	
MAGEL2, a candidate gene for autism and Prader-Willi syndrome	\$105,977	University of Alberta	
Longitudinal MRI Study of Brain Development in Fragile X	\$769,619	STANFORD UNIVERSITY	
Linking genetic mosaicism, neural circuit abnormalities and behavior	\$0	Brown University	
Linking circuit dynamics and behavior in a rat model of autism	\$0	University of California, San Francisco	
Language Development in Fragile X Syndrome	\$495,501	University of California, Davis	
Investigating the role of Tsc1 in neocortical circuit assembly	\$52,406	STANFORD UNIVERSITY	
Imaging of protein synthesis and ubiquitination in fragile x syndrome	\$195,000	Emory University	
Illuminating the role of glia in a zebrafish model of Rett syndrome	\$62,500	The Regents of the University of California, San Diego	
Identification of TSC cellular phenotypes using patient-derived iPSCs	\$193,750	Rutgers University	
Genotype-Phenotype Relationships in Fragile X Families	\$633,789	University of California, Davis	
Genetic Modifiers of Seizure Disorders in Fragile X Syndrome	\$261,539	Emory University	
Genetic and Developmental Analyses of Fragile X Mental Retardation Protein	\$383,322	Vanderbilt University	
Fragile X syndrome target analysis and its contribution to autism	\$124,725	Vanderbilt University	
FMRP regulates the pruning of cell-to-cell connections in the neocortex	\$79,500	UT SOUTHWESTERN MEDICAL CENTER	
FMRP and Pumilio co-regulate synaptogenesis by controlling Neuroglian expression	\$27,480	Vanderbilt University	
Emergence and Stability of Autism in Fragile X Syndrome	\$358,000	UNIVERSITY OF SOUTH CAROLINA	
Effects of Social Gaze Training on Brain and Behavior in Fragile X Syndrome	\$352,066	STANFORD UNIVERSITY	
Dysregulation of Protein Synthesis in Fragile X Syndrome and Other Developmental Disorders	\$1,221,847	National Institutes of Health	
Dysregulation of mTOR Signaling in Fragile X Syndrome	\$250,167	ALBERT EINSTEIN COLLEGE OF MEDICINE	
Dysregulation of mTOR Signaling in Fragile X Syndrome	\$164,833	ALBERT EINSTEIN COLLEGE OF MEDICINE	
Dysregulation of mTor/Tsc in 22q11DS Autism Model	\$62,500	GEORGE WASHINGTON UNIVERSITY	
Dysregulation of Mdm2-mediated p53 ubiquitination in autism mouse models	\$0	University of Illinois at Chicago	
Dissecting the 16p11.2 CNV endophenotype in induced pluripotent stem cells	\$54,400	University of California, San Francisco	
Development and afferent regulation of auditory neurons	\$376,200	Florida State University	
Dendritic 'translatome' in fragile X syndrome and autism	\$0	University of Michigan	
Cortico-striatal dysfunction in the eIF4E transgenic mouse model of autism	\$62,497	New York University	
Cortactin and Spine Dysfunction in Fragile X	\$33,763	University of California, Irvine	
Coordinate actions between methyl-CpG binding proteins in neuronal development	\$226,585	University of Wisconsin	

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Chloride homeostasis and GABA maturation in fragile X syndrome	\$231,750	Northwestern University	
Characterizing 22q11.2 abnormalities	\$62,498	Children's Hospital of Philadelphia	
Cell-type and circuit-specific functional deficits in cortex from gene disruptions linked to autism	\$30,000	University of North Carolina	
Calcium Channels as a Core Mechanism in the Neurobiology of ASD	\$35,000	Massachusetts General Hospital	
BDNF and the Restoration of Synaptic Plasticity in Fragile X and Autism	\$455,630	University of California, Irvine	
BAZ1B Haploinsufficiency and the Neuro-phenotypes of Williams Syndrome	\$59,000	The Regents of the University of California, Santa Barbara	
Autism phenotypes in Tuberous Sclerosis: Risk factors, features & architecture	\$0	King's College London	
Astrocytes contribution to tuberous sclerosis pathology	\$208,125	Yale University	
A Quantitative Study of Pyramidal Cells and Interneurons in the Cerebral Cortex	\$3,000	UNIVERSITY OF SOUTH CAROLINA	
A Novel Glial Specific Isoform of Cdkl5: Implications for the Pathology of Autism in Rett Syndrome	\$60,000	University of Nebraska	
A Novel Essential Gene for Human Cognitive Function	\$35,474	Harvard University	
Analysis of MEF2 in Cortical Connectivity and Autism-Associated Behaviors	\$56,042	McLean Hospital	
A mouse model for AUTS2-linked neurodevelopmental disorders	\$189,187	University of Illinois	
A Family-Genetic Study of Autism and Fragile X Syndrome	\$597,808	Northwestern University	
A Family-Genetic Study of Autism and Fragile X Syndrome	\$393,739	Northwestern University	
A cerebellar mutant for investigating mechanisms of autism in Tuberous Sclerosis	\$0	Boston Children's Hospital	
16p11.2 rearrangements: Genetic paradigms for neurodevelopmental disorders	\$100,000	University of Lausanne	